To the European Patent Office

Entry into the European phase (EPO as designated or elected Office)

European application number	EP04728561.4
PCT application number	PCT/EP2004/004245
PCT publication number	WO04101546
Applicant's or representative's reference	CJG/PB60227

1. Applicant

Particulars of the applicant(s) are contained in the international publication or were recorded by the International Bureau subsequent to the international publication. ☑

Changes which have not yet been recorded by the International Bureau are set out here: ☐

Address for correspondence

2. Representative 1

This is the representative who will be listed in the Register of European Patents and to whom notifications will be made

Name GODDARD, Carolyn, Janice

Address of place of business

CIP CN925.1
GlaxoSmithKline
980 Great West Road
Brentford, Middlesex TW8 9GS
United Kingdom

Telephone 0127 964 6051

Fax 020 8047 6894

e-mail

Any additional representative(s) is/are listed here: ☐

3. Authorisation

An individual authorisation is attached. ☐

A general authorisation has been registered under No: ☑ 31727

A general authorisation has been filed, but not yet registered. ☐

The authorisation filed with the EPO as PCT receiving Office expressly includes the European phase. ☐

4. Request for examination

Examination of the application under Art. 94 EPC is hereby requested. The examination fee is being (has been, will be) paid. ☑

Request for examination in an admissible non-EPO language: ☐

5. Copies

One or more additional sets of copies of the documents cited in the supplementary European search report are hereby requested. ☐

Number of additional sets of copies

6. Documents intended for proceedings before the EPO

6.1 Proceedings before the EPO as designated Office (PCT I) are to be based on the following documents:

the application documents published by the International Bureau (with all claims, description and drawings), where applicable with amended claims under Art. 19 PCT ☑

unless replaced by the amendments attached. ☐

Where necessary, clarifications should be attached as 'Other Documents'

6.2 Proceedings before the EPO as elected Office (PCT II) are to be based on the following documents:

the documents on which the international preliminary examination report is based, including any annexes ☐

unless replaced by the amendments attached. ☑

Where necessary, clarifications should be attached as 'Other Documents'

If the EPO as International Preliminary Examining Authority has been supplied with test reports, these may be used as the basis of proceedings before the EPO. ☐

7. Translations

Translations in one of the official languages of the EPO (English, French, German) are attached as crossed below:

- *In proceedings before the EPO as designated or elected Office (PCT I + II):*

Translation of the international application (description, claims, any text in the drawings) as originally filed, of the abstract as published and of any indication under Rule 13bis.3 and 13bis.4 PCT regarding biological material ☐

Translation of priority application(s) ☐

It is hereby declared that the international application as originally filed is a complete translation of the previous application (Rule 38(5) EPC) ☐

- *In addition, in proceedings before the EPO as designated Office (PCT I):*

Translation of amended claims and any statement under Art. 19 PCT, if the claims as amended are to form the basis for the proceedings before the EPO (see Section 6). ☐

- *In addition, in proceedings before the EPO as elected office (PCT II):*

Translation of annexes to the international preliminary examination report ☐

8. Biological material

The invention relates to and/or uses biological material deposited under Rule 28 EPC. ☐

The particulars referred to in Rule 28(1)(c) EPC (if not yet known, the depository institution and the identification reference(s) [number, symbols, etc.] of the depositor) are given in the international publication or in the translation submitted under Section 7 on: ☐

page(s) / line(s)

A copy of the receipt(s) of deposit issued by the depositary institution

is attached ☐

will be filed at a later date ☐

A waiver of the right to an undertaking from the requester pursuant to Rule 28(3) EPC is attached. ☐

9. Nucleotide and amino acid sequences

The items required under Rules 5.2 and 13ter PCT and Rule 111(3) EPC have already been furnished to the EPO. ☐

The sequence listing as part of the description is attached in PDF format. ☐

The sequence listing does not include matter that goes beyond the content of the application as filed. ☐

In addition, the sequence listing data is attached in computer-readable form in accordance with WIPO Standard 25. ☐

The sequence listing data in computer-readable form in accordance with WIPO Standard 25 is identical to the sequence listing in PDF format. ☐

10. Designation fees

10.1 It is currently intended to pay seven times the amount of the designation fee. The designation fees for all the EPC contracting states designated in the international application are thereby deemed to have been paid (Art. 2 No. 3 RFees). ☑

AT BE BG CH&LI CY CZ DE DK EE ES FI FR GB GR HU IE IS IT LT LU LV MC NL PL PT RO SE SI SK TR

10.2 The declaration in No. 10.1 does not apply. Instead, it is currently intended to pay fewer than seven designation fees for the following EPC contracting states designated in the international application: ☐

It is requested that no communications under Rule 108(3) EPC be issued in respect of any contracting states not indicated.

10.3 If an automatic debit order has been issued (Section 12), the EPO is authorised, on expiry of the basic period under Rule 107(1)(d) EPC, to debit seven times the amount of the designation fee. If states are indicated in No. 10.2, the EPO will debit designation fees for those states only, unless instructed otherwise before the basic period expires. ☑

11. Extension of the European patent

This application is also considered as being a request for extension to all the non-contracting states to the EPC designated in the international application with which "extension agreements" were in force on the date of filing the international application. However, the extension only takes effect if the prescribed extension fee is paid. ☑

It is currently intended to pay the extension fee for the following states:

HR LV

12. Debit from deposit account ☑

Currency: EUR

The European Patent Office is hereby authorised to debit from the following deposit account any fees and costs indicated on the fees page.

Deposit account number: 28050015
Account holder: GlaxoSmithKline

13. Reimbursements (if any) should be made to the following EPO deposit account: ☑

Number and account holder: 28050015, GlaxoSmithKline

14. Fees

		Factor applied	Fee schedule	Amount to be paid
14-1	005 Designation fee	7	75.00	525.00
14-2	006 Examination fee	1 (50%)	1 430.00	715.00
14-3	015 Claims fee	17	40.00	680.00
14-4	020 Basic national fee for an international application	1	90.00	90.00
14-5	402 Extension fee LT	1	102.00	102.00
14-6	403 Extension fee LV	1	102.00	102.00
14-7	407 Extension fee for Croatia	1	102.00	102.00
	Total:		**EUR**	**2 316.00**

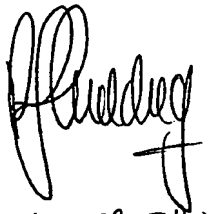
15. Annotations

15-1. Note (for EPO) (EP Phase)

Extension State (LT) (GODDARD, Carolyn Janice; 18/10/2005)
Please note Lithuania should be designated as an extension state - there is no provision for this on the electronic form. Extension fee has been paid.

16. Signature(s) of applicant(s) or representative

Place: **Brentford**
Date: **19 October 2005**
Signed by: **/F Reardon/**



PETER JOHN GIDDINGS

GA 31727

Capacity: **GlaxoSmithKline (Applicant, as employee)**

For employees (Art. 133(3) EPC) having a general authorisation:
General authorisation No.